HAND SANITIZING WIPES- alcohol cloth Trinity Packaging Supply Limited Liability Company

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

This is a hand sanitizer manufactured according to the Temporary Policy for Preparation of Certain Alcohol-Based Hand Sanitizer Products During the Public Health Emergency (CoViD-19); Guidance for Industry.

The hand sanitizer is manufactured using only the following United States Pharmacopoeia (USP) grade ingredients in the preparation of the product (percentage in final product formulation) consistent with World Health Organization (WHO) recommendations:

- a. Alcohol (ethanol) (USP or Food Chemical Codex (FCC) grade) (75%, volume/volume (v/v)) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.
- b. Glycerol (1.45% v/v).
- c. Hydrogen peroxide (0.125% v/v).
- d. Sterile distilled water or boiled cold water.

The firm does not add other active or inactive ingredients. Different or additional ingredients may impact the quality and potency of the product.

Active Ingredient(s)

Alcohol 75% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizing Wipes

Use

Hand Sanitizing Wipes help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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away.

Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

purified water USP

Package Label - Principal Display Panel



373 mL NDC: 75665-200-10

HAND SANITIZING WII	PES						
Product Information							
Product Type Route of Administration	HUMAN OTC DRUG	Item Code	(Source)	NDC:75665-200			
Active Ingredient/Active Moiety							
Ingredient Name			Basis of Strengt	h Strength			

ALCOHOL	(UNII: 3K9958V90M)	(ALCOHOL - UNII:3K9958V90M)
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ALCOHOL

Inactive Ingredients							
Ingredient Name				Strength			
W	WATER (UNII: 059QF0KO0R)						
Packaging							
#	Item Code	Package Description	Marketing Star	rt Date Marketing End Date			
1	NDC:75665-200-10	373 mL in 1 POUCH; Type 0: Not a Combination Product	03/30/2020				
Marketing Information							
	Marketing Categor	y Application Number or Monograph Citation	Marketing Sta	rt Date Marketing End Date			
0	TC monograph not fin	al part333A	03/30/2020				

Labeler - Trinity Packaging Supply Limited Liability Company (042641690)

Establishment

Name	Address	ID/FEI	Business Operations
Trinity Packaging Supply Limited Liability Company		042641690	manufacture(75665-200)

Revised: 5/2020

Trinity Packaging Supply Limited Liability Company